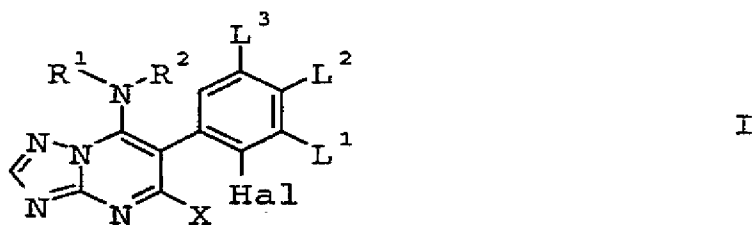


Amendments to the CLAIMS

1. (Withdrawn) A substituted 6-(2-halogenphenyl)-triazolopyrimidine of formula I



in which

Hal is halogen;

L¹, L³ independently denote hydrogen, halogen, or C₁-C₄-alkyl;

L² is hydrogen, halogen, C₁-C₄-haloalkyl, or NH₂, NHR^b, or N(R^b)₂,

R^b is C₁-C₈-alkyl, C₃-C₁₀-alkenyl, C₃-C₁₀-alkynyl, C₁-C₆-haloalkyl, C₃-C₆-haloalkenyl, C₃-C₆-haloalkynyl, C₁-C₈-alkoxy-C₁-C₈-alkyl, C₁-C₈-alkylthio-C₁-C₈-alkyl, C₃-C₁₀-cycloalkyl, or C(=O)-A, in which

A is hydrogen, hydroxy, C₁-C₈-alkyl, C₁-C₈-alkoxy, C₁-C₆-halogenalkoxy, C₁-C₈-alkylamino or di-(C₁-C₈-alkyl)amino;

wherein at least one from L¹, L², and L³ is not hydrogen;

X is halogen, cyano, C₁-C₆-alkyl, C₁-C₆-alkoxy, C₁-C₆-haloalkoxy or C₃-C₈-alkenyloxy;

R^1 denote C_1 - C_{10} -alkyl, C_2 - C_{10} -alkenyl, C_2 - C_{10} -alkynyl, or C_4 - C_{10} -alkadienyl, C_2 - C_{10} -haloalkenyl

wherein R^1 may be unsubstituted or may carry one to three groups R^a ,

R^a is cyano, nitro, hydroxyl, C_1 - C_6 -alkyl, C_3 - C_6 -cycloalkyl, C_1 - C_6 -alkoxy, C_1 - C_6 -alkylthio, C_1 - C_6 -alkylamino, di- C_1 - C_6 -alkylamino, C_2 - C_6 -alkenyl, C_2 - C_6 -alkenyloxy, C_2 - C_6 -alkynyl, C_3 - C_6 -alkynyloxy, or C_1 - C_4 -alkylenedioxy;

and R^2 is hydrogen.

2. (Withdrawn, Currently amended) The ~~compound~~ substituted 6-(2-halogenphenyl)-triazolopyrimidine of formula I according to claim 1, in which

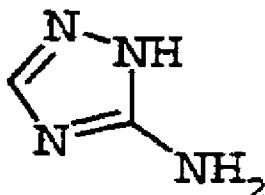
R^1 is straight chained or branched C_2 - C_6 -alkenyl, or a straight chained or branched C_1 - C_6 -alkyl.

3. (Withdrawn, Currently amended) The ~~compound~~ substituted 6-(2-halogenphenyl)-triazolopyrimidine of formula I according to claim 1 or 2 in which X is halogen.

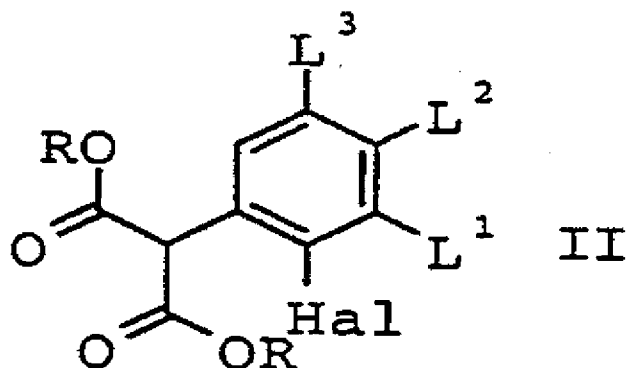
4. (Withdrawn, Currently amended) The ~~compound~~ substituted 6-(2-halogenphenyl)-triazolopyrimidine of formula I according to claim 1 in which the 6-(2-halogenphenyl)group represents one of the following moieties:

2,3,5-trifluorophenyl; 2-F,4-CF₃-phenyl; 2-F,5-CH₃-phenyl; 2-Cl,4-F-phenyl; 2-F,4-Cl-phenyl; 2-F,4-Br-phenyl; 2-Cl,4-Br-phenyl; 2,3-difluorophenyl; 2,4-difluorophenyl; 2,4,5-trifluorophenyl; 2,3,4-trifluorophenyl; 2-F,4-NHC(O)CH₃-phenyl; 2-Br,3,5-difluorophenyl; 2-F,4-NO₂-phenyl; and 2-Cl,4-NO₂-phenyl.

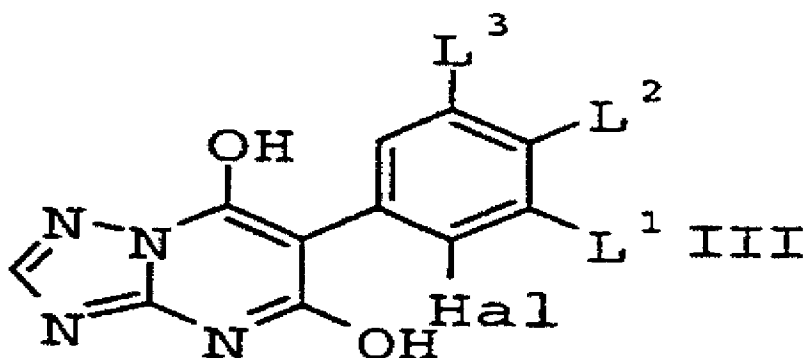
5. (Withdrawn, Currently amended) A process for the preparation of compounds the substituted 6-(2-halogenphenyl)-triazolopyrimidine of formula I as defined in claim 3 which comprises reacting 5-amino-1,2,4-triazole



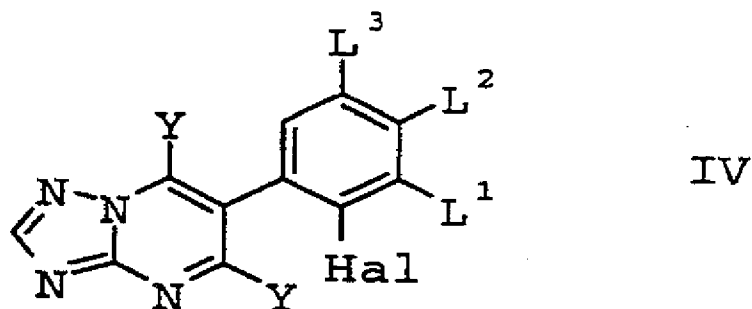
with 2-phenyl-substituted malonic acid ester of formula II,



wherein Hal, L¹, L², and L³ are as defined in formula I, and R denotes C₁-C₆-alkyl, under alkaline conditions, to yield compounds of formula III,



which are subsequently treated with a halogenating agent to give 5,7-dihalogen-6-phenyl-triazolopyrimidines of formula IV

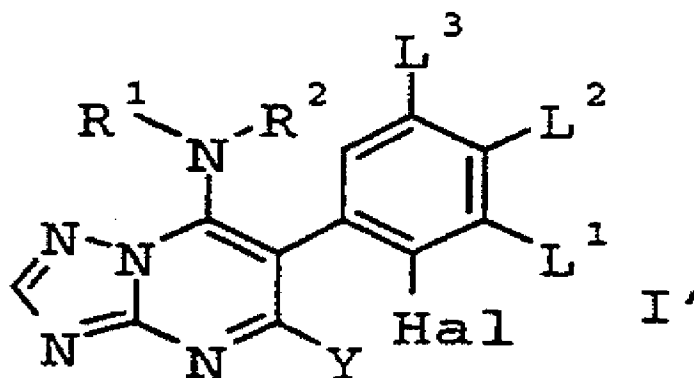


in which Y is halogen, and which is reacted with an amine of formula V



in which R^1 and R^2 are as defined in claim 1 to produce compounds of formula I, as defined in claim 1.

6. (Withdrawn, Currently amended) A process for the preparation of compounds the substituted 6-(2-halogenphenyl)-triazolopyrimidine of formula I according to claim 1 wherein X is cyano, C_1 - C_{10} -alkoxy, or C_1 - C_6 -haloalkoxy, which comprises reacting 5-halogen-triazolopyrimidine of formula I',



wherein Y is halogen, with compounds of formula VI,

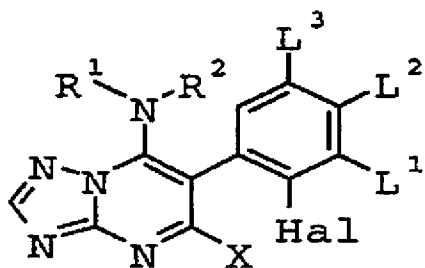


which are, dependent from the value of X' to be introduced, an inorganic cyano salt, an alkoxylate, haloalkoxylate or an alkenyloxylate, respectively, wherein M is ammonium-, tetraalkylammonium-, alkalimetal- or alkaline earth metal cation, to produce compounds of formula I.

7. (Withdrawn) An intermediate of formulae II, III, or IV as defined in claim 5, in which the 6-(2-halogenphenyl)group represents one of the following moieties:

2,3,5-trifluorophenyl; 2-F,4-CF₃-phenyl; 2-F,5-CH₃-phenyl; 2-Cl,4-F-phenyl; 2-F,4-Cl-phenyl; 2-F,4-Br-phenyl; 2-Cl,4-Br-phenyl; 2,3-difluorophenyl; 2,4,5-trifluorophenyl; 2,3,4-trifluorophenyl; 2-F,4-NHC(O)CH₃-phenyl; 2-Br,3,5-difluorophenyl; 2-F,4-NO₂-phenyl; and 2-Cl,4-NO₂-phenyl.

8. (Withdrawn, Currently amended) A composition suitable for controlling phytopathogenic fungi, comprising a solid or liquid carrier and a compound the substituted 6-(2-halogenphenyl)-triazolopyrimidine of the formula I as claimed in claim 1.
9. (Withdrawn, Currently amended) A method for controlling phytopathogenic fungi, which comprises treating the fungi or the materials, plants, the soil or the seed to be protected against fungal attack with an effective amount of ~~a compound~~ the substituted 6-(2-halogenphenyl)-triazolopyrimidine of the formula I as claimed in claim 1.
10. (Previously presented) A substituted 6-(2-halogenphenyl)-triazolopyrimidine of formula I



I

in which

Hal is halogen;

L¹, L³ independently denote hydrogen, halogen, or C₁-C₄-alkyl;

L^2 is hydrogen, halogen, C_1 - C_4 -haloalkyl, or NH_2 , NHR^b , or $N(R^b)_2$,

R^b is C_1 - C_8 -alkyl, or $C(=O)$ -A, in which

A is C_1 - C_8 -alkyl;

wherein at least one from L^1 , L^2 , and L^3 is not hydrogen;

X is halogen, C_1 - C_6 -alkyl, or C_1 - C_6 -alkoxy;

R^1 and R^2 together with the interjacent nitrogen atom represent a saturated or partially unsaturated 5- or 6-membered heterocycle, containing one nitrogen atom or one nitrogen atom and one sulfur atom, which ring may be substituted by one to three R^a radicals;

R^a is C_1 - C_6 alkyl.

11. (Currently amended) The compound substituted 6-(2-halogenphenyl)-triazolopyrimidine of formula I according to claim 10, in which

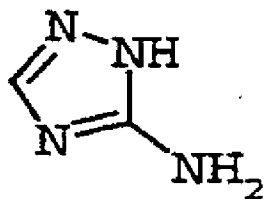
R^1 and R^2 together with the interjacent nitrogen atom represent a saturated or partially unsaturated 5- or 6-membered heterocycle, containing one nitrogen atom or one nitrogen atom and one sulfur atom, being optionally substituted with one or two C_1 - C_4 -alkyl groups.

12. (Currently amended) The compound substituted 6-(2-halogenphenyl)-triazolopyrimidine of formula I according to claim 10 in which R^1 and R^2 together with the interjacent nitrogen atom represent a saturated or partially unsaturated 5- or 6-membered heterocycle, containing one nitrogen atom or one nitrogen atom and one sulfur atom, being optionally

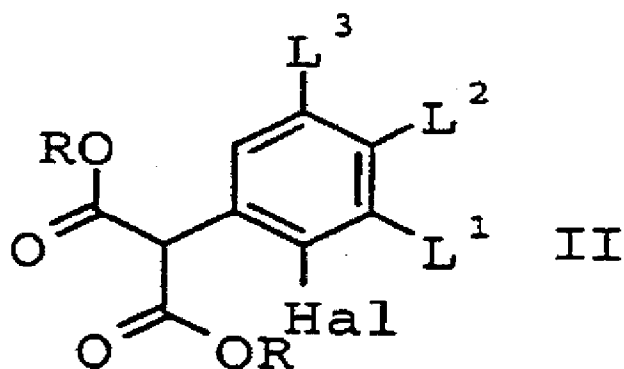
substituted with one or two methyl groups.

13. (Currently amended) The ~~compound~~ substituted 6-(2-halogenphenyl)-triazolopyrimidine of formula I according to claim 10 in which X is halogen.
14. (Currently amended) The ~~compound~~ substituted 6-(2-halogenphenyl)-triazolopyrimidine of formula I according to claim 10 in which the 6-(2-halogenphenyl)group represents one of the following moieties:

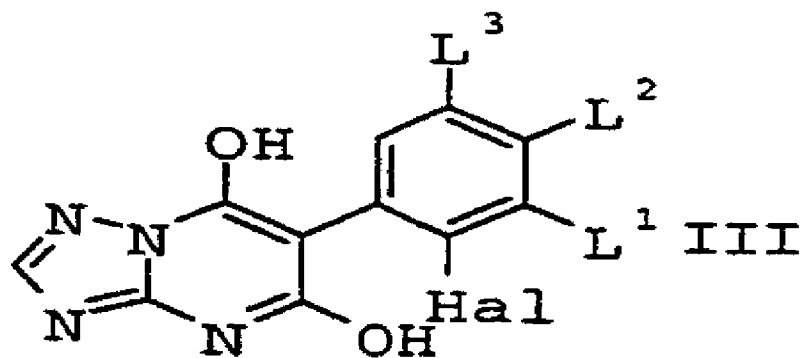
2,3,5-trifluorophenyl; 2-F,4-CF₃-phenyl; 2-F,5-CH₃-phenyl; 2-Cl,4-F-phenyl; 2-F,4-Cl-phenyl; 2-F,4-Br-phenyl; 2-Cl,4-Br-phenyl; 2,3-difluorophenyl; 2,4-difluorophenyl; 2,4,5-trifluorophenyl; 2,3,4-trifluorophenyl; 2-F,4-NHC(O)CH₃-phenyl; and 2-Br,3,5-difluorophenyl.
15. (Withdrawn, Currently amended) A process for the preparation of the ~~compound~~ substituted 6-(2-halogenphenyl)-triazolopyrimidine of formula I as defined in claim 13 which comprises reacting 5-amino-1,2,4-triazole



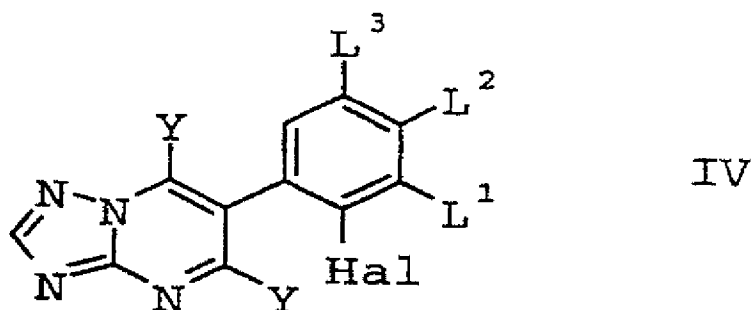
with 2-phenyl-substituted malonic acid ester of formula II,



wherein Hal, L¹, L², and L³ are as defined in formula I, and R denotes C₁-C₆-alkyl, under alkaline conditions, to yield compounds of formula III,



which are subsequently treated with a halogenating agent to give 5,7-dihalogen-6-phenyl-triazolopyrimidines of formula IV

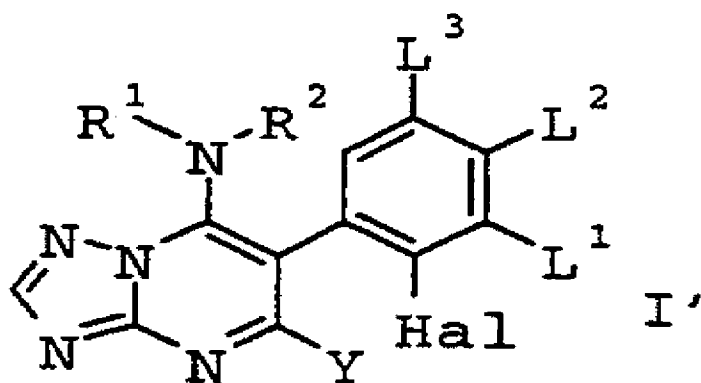


in which Y is halogen, and which is reacted with an amine of formula V



in which R¹ and R² are as defined in claim 10 to produce compounds of formula I, as defined in claim 13.

16. (Withdrawn, Currently amended) A process for the preparation the compound substituted 6-(2-halogenphenyl)-triazolopyrimidine of formula I according to claim 10 wherein X is C₁-C₁₀-alkoxy, which comprises reacting 5-halogen-triazolopyrimidine of formula I',



wherein Y is halogen, with compounds of formula VI,



which is an alkoxylate, wherein M is ammonium-, tetraalkylammonium-, alkalimetal- or alkaline earth metal cation, to produce compounds of formula I.

17. (Currently amended) A composition suitable for controlling phytopathogenic fungi, comprising a solid or liquid carrier and ~~a compound~~ the substituted 6-(2-halogenphenyl)-triazolopyrimidine of the formula I as claimed in claim 10.
18. (Withdrawn, Currently amended) A method for controlling phytopathogenic fungi, which comprises treating the fungi or the materials, plants, the soil or the seed to be protected against fungal attack with an effective amount of ~~a compound~~ the substituted 6-(2-halogenphenyl)-triazolopyrimidine of the formula I as claimed in claim 10.